

**Chemistry
Higher level
Paper 2**

Monday 14 November 2016 (morning)

Candidate session number

2 hours 15 minutes

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Instructions to candidates

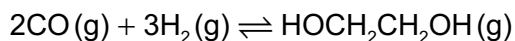
- Write your session number in the boxes above.
- Do not open this examination paper until instructed to do so.
- Answer all questions.
- Write your answers in the boxes provided.
- A calculator is required for this paper.
- A clean copy of the **chemistry data booklet** is required for this paper.
- The maximum mark for this examination paper is **[95 marks]**.



Answer **all** questions. Write your answers in the boxes provided.

1. Ethane-1,2-diol, $\text{HOCH}_2\text{CH}_2\text{OH}$, has a wide variety of uses including the removal of ice from aircraft and heat transfer in a solar cell.

(a) Ethane-1,2-diol can be formed according to the following reaction.



(i) Deduce the equilibrium constant expression, K_c , for this reaction.

[1]

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(ii) State how increasing the pressure of the reaction mixture at constant temperature will affect the position of equilibrium and the value of K_c .

[2]

Position of equilibrium:

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K_c :

.....

(iii) Calculate the enthalpy change, ΔH^\ominus , in kJ, for this reaction using section 11 of the data booklet. The bond enthalpy of the carbon–oxygen bond in CO(g) is 1077 kJ mol^{-1} .

[3]

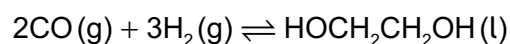
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(Question 1 continued)

- (b) (i) Calculate ΔH^\ominus , in kJ, for this similar reaction below using ΔH_f^\ominus data from section 12 of the data booklet. ΔH_f^\ominus of $\text{HOCH}_2\text{CH}_2\text{OH}(\text{l})$ is $-454.8 \text{ kJ mol}^{-1}$. [1]



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- (ii) Deduce why the answers to (a)(iii) and (b)(i) differ. [1]

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- (iii) ΔS^\ominus for the reaction in (b)(i) is -620.1 J K^{-1} . Comment on the decrease in entropy. [1]

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- (iv) Calculate the value of ΔG^\ominus , in kJ, for this reaction at 298 K using your answer to (b)(i). (If you did not obtain an answer to (b)(i), use -244.0 kJ , but this is not the correct value.) [2]

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(Question 1 continued)

- (v) Comment on the statement that the reaction becomes less spontaneous as temperature is increased.

[1]

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- (c) Determine the average oxidation state of carbon in ethene and in ethane-1,2-diol.

[2]

Ethene:

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Ethane-1,2-diol:

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- (d) Explain why the boiling point of ethane-1,2-diol is significantly greater than that of ethene.

[2]

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- (e) Ethane-1,2-diol can be oxidized first to ethanedioic acid, $(\text{COOH})_2$, and then to carbon dioxide and water. Suggest the reagents needed to oxidize ethane-1,2-diol.

[1]

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(This question continues on the following page)



(Question 1 continued)

- (f) Predict the ^1H NMR data for ethanedioic acid and ethane-1,2-diol by completing the table.

[2]

	Number of signals	Splitting pattern
Ethanedioic acid:
Ethane-1,2-diol:	Not required



2. The concentration of a solution of a weak acid, such as ethanedioic acid, can be determined by titration with a standard solution of sodium hydroxide, NaOH(aq).

(a) Distinguish between a weak acid and a strong acid.

[1]

Weak acid:

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Strong acid:

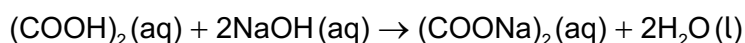
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(b) Suggest why it is more convenient to express acidity using the pH scale instead of using the concentration of hydrogen ions.

[1]

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- (c) 5.00 g of an impure sample of hydrated ethanedioic acid, $(\text{COOH})_2 \cdot 2\text{H}_2\text{O}$, was dissolved in water to make 1.00 dm^3 of solution. 25.0 cm^3 samples of this solution were titrated against a $0.100 \text{ mol dm}^{-3}$ solution of sodium hydroxide using a suitable indicator.



The mean value of the titre was 14.0 cm^3 .

(i) Suggest a suitable indicator for this titration. Use section 22 of the data booklet.

[1]

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(This question continues on the following page)



(Question 2 continued)

(ii) Calculate the amount, in mol, of NaOH in 14.0 cm^3 of 0.100 mol dm^{-3} solution. [1]

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(iii) Calculate the amount, in mol, of ethanedioic acid in each 25.0 cm³ sample. [1]

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(iv) Determine the percentage purity of the hydrated ethanedioic acid sample. [3]

[illegible]

(d) Draw the Lewis (electron dot) structure of the ethanedioate ion, OOC^-COO^- . [1]

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(This question continues on the following page)



(Question 2 continued)

- (e) Outline why all the C–O bond lengths in the ethanedioate ion are the same length and suggest a value for them. Use section 10 of the data booklet.

[2]

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- (f) Explain how ethanedioate ions act as ligands.

[2]

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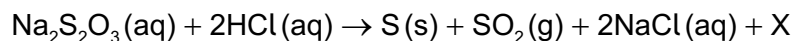
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3. Sodium thiosulfate solution reacts with dilute hydrochloric acid to form a precipitate of sulfur at room temperature.



- (a) Identify the formula and state symbol of X. [1]

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- (b) Suggest why the experiment should be carried out in a fume hood or in a well-ventilated laboratory. [1]

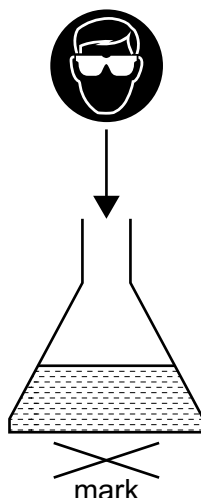
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(Question 3 continued)

- (c) The precipitate of sulfur makes the mixture cloudy, so a mark underneath the reaction mixture becomes invisible with time.



10.0 cm³ of 2.00 mol dm⁻³ hydrochloric acid was added to a 50.0 cm³ solution of sodium thiosulfate at temperature, T₁. Students measured the time taken for the mark to be no longer visible to the naked eye. The experiment was repeated at different concentrations of sodium thiosulfate.

Experiment	[Na ₂ S ₂ O ₃ (aq)] / mol dm ⁻³	Time, t, for mark to disappear / s ± 1 s	$\frac{1}{t} / 10^{-3} \text{ s}^{-1}$
1	0.150	23	43.5
2	0.120	27	37.0
3	0.090	36	27.8
4	0.060	60	16.7
5	0.030	111	9.0

* The reciprocal of the time in seconds can be used as a measure of the rate of reaction.

[Source: Adapted from <http://www.flinnsci.com/>]

Show that the hydrochloric acid added to the flask in experiment 1 is in excess.

[2]

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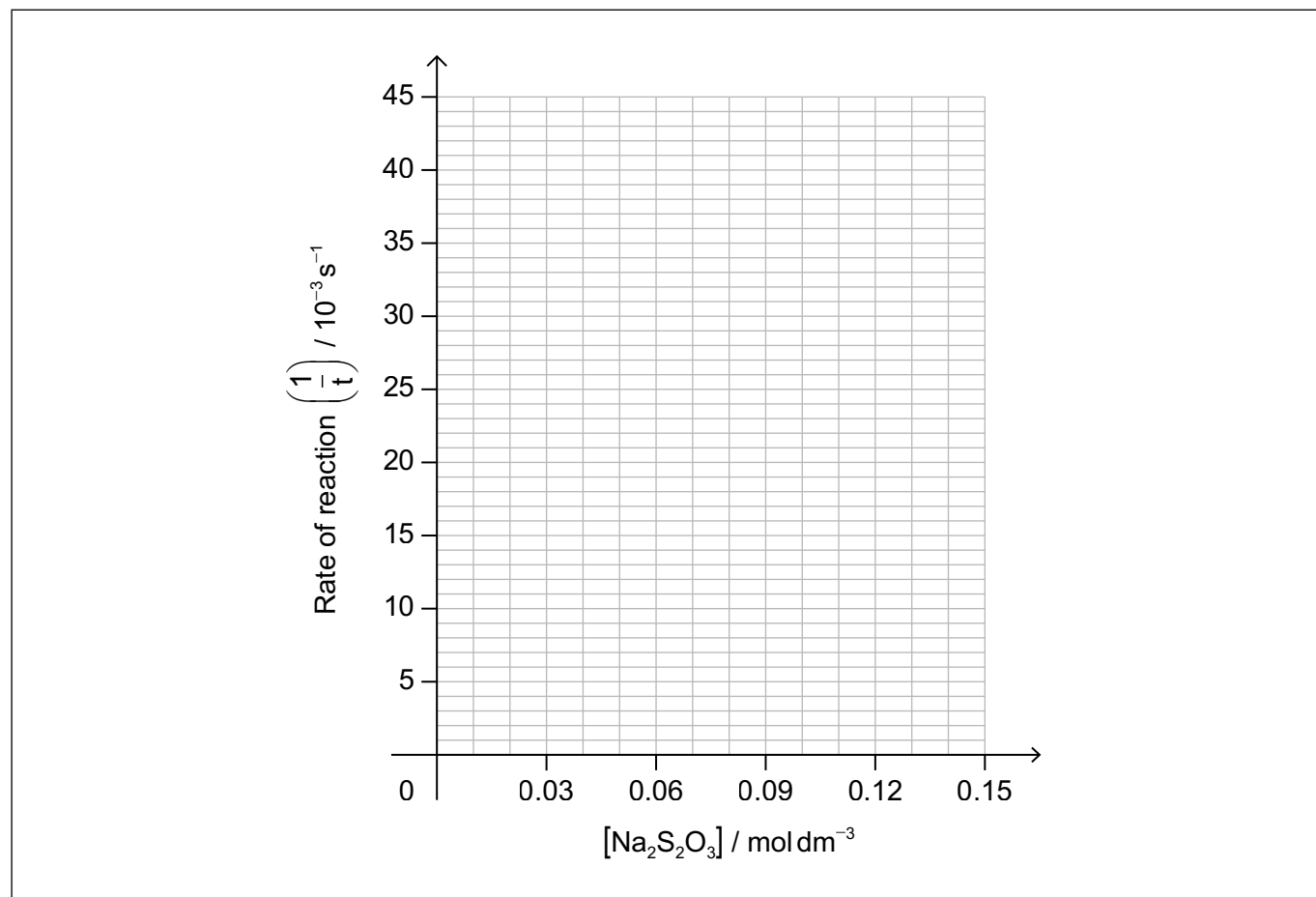
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(Question 3 continued)

- (d) Draw the best fit line of $\frac{1}{t}$ against concentration of sodium thiosulfate on the axes provided. [2]



- (e) (i) Using the graph, explain the order of reaction with respect to sodium thiosulfate. [2]

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(Question 3 continued)

- (ii) In a different experiment, this reaction was found to be first order with respect to hydrochloric acid. Deduce the overall rate expression for the reaction.

[1]

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- (f) A student decided to carry out another experiment using $0.075 \text{ mol dm}^{-3}$ solution of sodium thiosulfate under the same conditions. Determine the time taken for the mark to be no longer visible.

[2]

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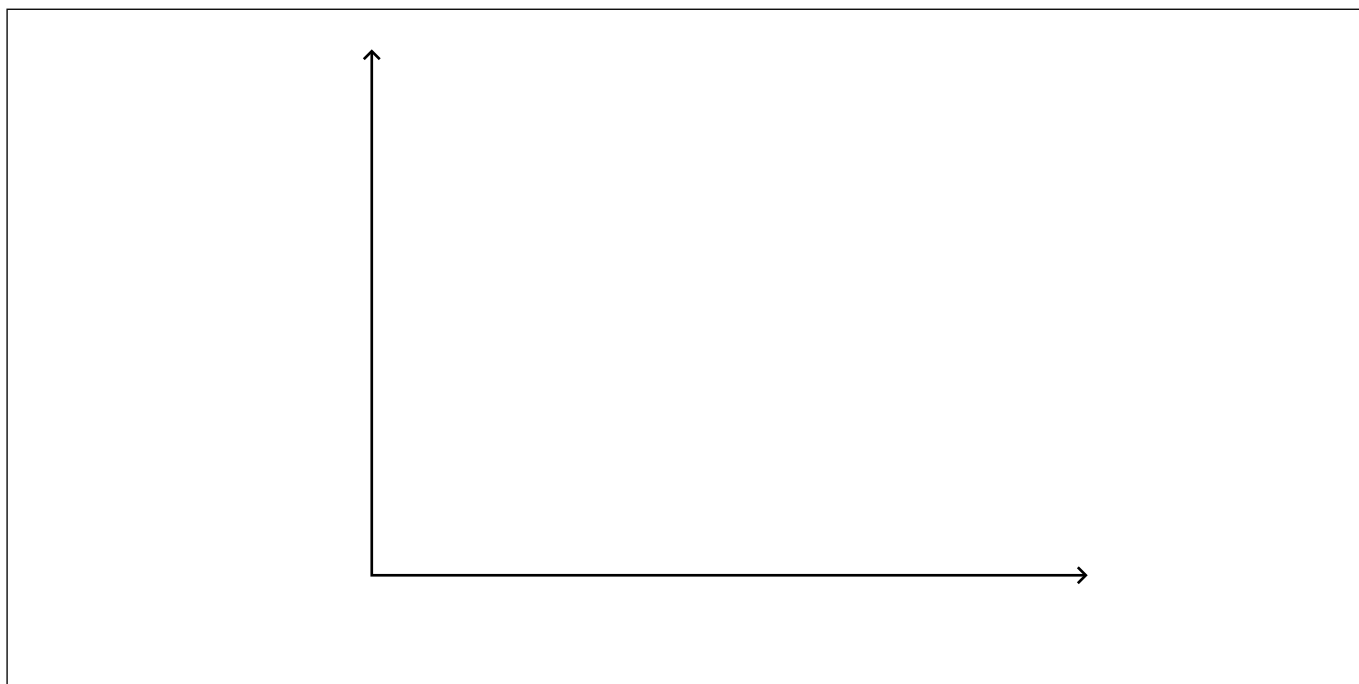
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- (g) An additional experiment was carried out at a higher temperature, T_2 .

- (i) On the same axes, sketch Maxwell–Boltzmann energy distribution curves at the two temperatures T_1 and T_2 , where $T_2 > T_1$.

[2]



(This question continues on the following page)



(Question 3 continued)

- (ii) Explain why a higher temperature causes the rate of reaction to increase. [2]

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- (h) Suggest one reason why the values of rates of reactions obtained at higher temperatures may be less accurate. [1]

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4. Magnesium is a group 2 metal which exists as a number of isotopes and forms many compounds.

(a) State the nuclear symbol notation, A_ZX , for magnesium-26.

[1]

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(b) Mass spectroscopic analysis of a sample of magnesium gave the following results:

	% abundance
Mg-24	78.60
Mg-25	10.11
Mg-26	11.29

Calculate the relative atomic mass, A_r , of this sample of magnesium to two decimal places.

[2]

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(c) Magnesium ions produce no emission or absorption lines in the visible region of the electromagnetic spectrum. Suggest why most magnesium compounds tested in a school laboratory show traces of yellow in the flame.

[1]

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(This question continues on the following page)



(Question 4 continued)

- (d) (i) Explain the convergence of lines in a hydrogen emission spectrum. [1]

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- (ii) State what can be determined from the frequency of the convergence limit. [1]

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- (e) Magnesium burns in air to form a white compound, magnesium oxide. Formulate an equation for the reaction of magnesium oxide with water. [1]

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- (f) Describe the trend in acid-base properties of the oxides of period 3, sodium to chlorine. [2]

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- (g) In addition to magnesium oxide, magnesium forms another compound when burned in air. Suggest the formula of this compound. [1]

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(This question continues on the following page)



(Question 4 continued)

- (h) Describe the structure and bonding in solid magnesium oxide.

[2]

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- (i) Magnesium chloride can be electrolysed.

- (i) Deduce the half-equations for the reactions at each electrode when **molten** magnesium chloride is electrolysed, showing the state symbols of the products. The melting points of magnesium and magnesium chloride are 922 K and 987 K respectively.

[2]

Anode (positive electrode):

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Cathode (negative electrode):

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- (ii) Identify the type of reaction occurring at the cathode (negative electrode).

[1]

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- (iii) State the products when a very **dilute** aqueous solution of magnesium chloride is electrolysed.

[2]

Anode (positive electrode):

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Cathode (negative electrode):

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(This question continues on the following page)



(Question 4 continued)

- (j) Standard electrode potentials are measured relative to the standard hydrogen electrode. Describe a standard hydrogen electrode.

[2]

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- (k) A magnesium half-cell, $\text{Mg(s)}/\text{Mg}^{2+}(\text{aq})$, can be connected to a copper half-cell, $\text{Cu(s)}/\text{Cu}^{2+}(\text{aq})$.

- (i) Formulate an equation for the spontaneous reaction that occurs when the circuit is completed.

[1]

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- (ii) Determine the standard cell potential, in V, for the cell. Refer to section 24 of the data booklet.

[1]

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- (iii) Predict, giving a reason, the change in cell potential when the concentration of copper ions increases.

[2]

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5. Propane and propene are members of different homologous series.

(a) Draw the full structural formulas of propane and propene.

[1]

Propane:

Propene:

(b) (i) Draw diagrams to show how sigma (σ) and pi (π) bonds are formed between atoms.

[2]

Sigma (σ):

Pi (π):

(This question continues on the following page)



(Question 5 continued)

- (ii) State the number of sigma (σ) and pi (π) bonds in propane and propene. [2]

	Number of sigma (σ) bonds	Number of pi (π) bonds
Propane
Propene

- (c) Both propane and propene react with bromine.

- (i) State an equation and the condition required for the reaction of 1 mol of propane with 1 mol of bromine. [2]

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- (ii) State an equation for the reaction of 1 mol of propene with 1 mol of bromine. [1]

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- (iii) State the type of each reaction with bromine. [1]

Propane:

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Propene:

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(This question continues on the following page)



(Question 5 continued)

- (d) Construct the mechanism of the formation of 2-bromopropane from hydrogen bromide and propene using curly arrows to denote the movement of electrons.

[3]



6. One structural isomer of C_4H_9Br is a chiral molecule.

- (a) Draw the three-dimensional shape of each enantiomer of this isomer showing their spatial relationship to each other. [2]

- (b) When one enantiomer undergoes substitution by alkaline hydrolysis approximately 75 % of the product molecules show inversion of configuration. Comment on the mechanisms that occur. [2]

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- (c) Suggest why the rate of alkaline hydrolysis of an enantiomer of iodopropane is greater than that of an enantiomer of bromopropane. [1]

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7. This question is about the weak acid methanoic acid, HCOOH .

- (a) Calculate the pH of $0.0100 \text{ mol dm}^{-3}$ methanoic acid stating any assumption you make.
 $K_a = 1.6 \times 10^{-4}$.

[3]

Calculation:

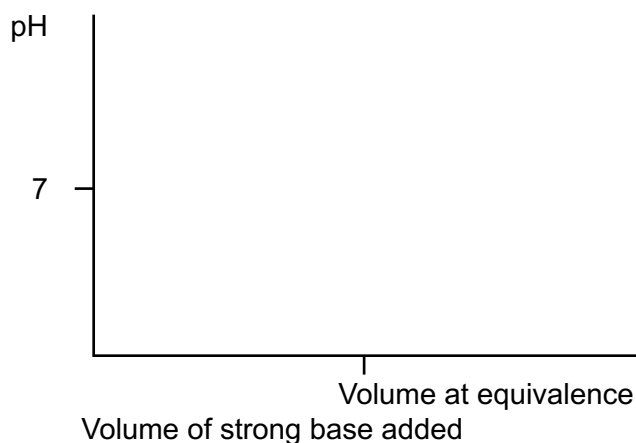
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Assumption:

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- (b) (i) Sketch a graph of pH against volume of a strong base added to a weak acid showing how you would determine pK_a for the weak acid.

[2]



(This question continues on the following page)



(Question 7 continued)

- (ii) Explain, using an equation, why the pH increases very little in the buffer region when a small amount of alkali is added.

[2]

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Answers written on this page
will not be marked.



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